

GenCore version 5.1.3
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OM nucleic - nucleic search, using SW model

Run on: October 6, 2002, 12:40:30 : Search time 284 Seconds
(without alignments)
120.909 Million cell updates/sec

Title: US-09-754-468-47
Perfect score: 20
Sequence: 1 gattagcataataaattc 20

Scoring table: IDENTITY_NUC
Gapop 10.0, Gapexcl 1.0

Searched: 1736436 seqs, 858457221 residues
Total number of hits satisfying chosen parameters: 3472872

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : N_Genesec_032802.1
1: /net/abs06/SIDSL1/gcgdata/hold-genesec/genesecq-emb1/NA1980.DAT:*
2: /net/abs06/SIDSL1/gcgdata/hold-genesec/genesecq-emb1/NA1981.DAT:*
3: /net/abs06/SIDSL1/gcgdata/hold-genesec/genesecq-emb1/NA1982.DAT:*
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5: /net/abs06/SIDSL1/gcgdata/hold-genesec/genesecq-emb1/NA1984.DAT:*
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8: /net/abs06/SIDSL1/gcgdata/hold-genesec/genesecq-emb1/NA1987.DAT:*
9: /net/abs06/SIDSL1/gcgdata/hold-genesec/genesecq-emb1/NA1988.DAT:*
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19: /net/abs06/SIDSL1/gcgdata/hold-genesec/genesecq-emb1/NA1998.DAT:*
20: /net/abs06/SIDSL1/gcgdata/hold-genesec/genesecq-emb1/NA1999.DAT:*
21: /net/abs06/SIDSL1/gcgdata/hold-genesec/genesecq-emb1/NA2000.DAT:*
22: /net/abs06/SIDSL1/gcgdata/hold-genesec/genesecq-emb1/NA2001A.DAT:*
23: /net/abs06/SIDSL1/gcgdata/hold-genesec/genesecq-emb1/NA2001B.DAT:*
24: /net/abs06/SIDSL1/gcgdata/hold-genesec/genesecq-emb1/NA2002.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length DB	Hit	Description
1	20	100.0	836	22	AAH25706
2	20	100.0	836	22	AAH25707
3	20	100.0	2754	20	AAH38420
4	20	100.0	3811	20	AAH38421
5	18.4	92.0	325	21	AAH38422
6	18.4	92.0	645	22	AAH38423
7	18.4	92.0	1972	22	AAH38424
8	18.4	92.0	1972	22	AAH38425
9	18.4	92.0	1976	22	AAH38426

10	17.4	87.0	3153	21	AAH38427	Human secreted pro
11	16.4	82.0	401	22	AAH38428	Human neutrophil
12	16.4	82.0	401	22	AAH38429	Human neutrophil
13	16.4	82.0	401	22	AAH38430	Human neutrophil
14	16.4	82.0	401	22	AAH38431	Human neutrophil
15	16.4	82.0	401	22	AAH38432	Human neutrophil
16	16.4	82.0	401	22	AAH38433	Human neutrophil
17	16.4	82.0	401	22	AAH38434	Human neutrophil
18	16.4	82.0	401	22	AAH38435	Human neutrophil
19	16.4	82.0	401	22	AAH38436	Human neutrophil
20	16.4	82.0	401	22	AAH38437	Human neutrophil
21	16.4	82.0	401	22	AAH38438	Human neutrophil
22	16.4	82.0	401	22	AAH38439	Human neutrophil
23	16.4	82.0	401	22	AAH38440	Human neutrophil
24	16.4	82.0	401	22	AAH38441	Human neutrophil
25	16.4	82.0	401	22	AAH38442	Human neutrophil
26	16.4	82.0	401	22	AAH38443	Human neutrophil
27	16.4	82.0	401	22	AAH38444	Human neutrophil
28	16.4	82.0	401	22	AAH38445	Human neutrophil
29	16.4	82.0	401	22	AAH38446	Human neutrophil
30	16.4	82.0	401	22	AAH38447	Human neutrophil
31	16.4	82.0	401	22	AAH38448	Human neutrophil
32	16.4	82.0	401	22	AAH38449	Human neutrophil
33	16.4	82.0	401	22	AAH38450	Human neutrophil
34	16.4	82.0	401	22	AAH38451	Human neutrophil
35	16.4	82.0	401	22	AAH38452	Human neutrophil
36	16.4	82.0	401	22	AAH38453	Human neutrophil
37	16.4	82.0	401	22	AAH38454	Human neutrophil
38	16.4	82.0	401	22	AAH38455	Human neutrophil
39	16.4	82.0	401	22	AAH38456	Human neutrophil
40	16.4	82.0	401	22	AAH38457	Human neutrophil
41	16.4	82.0	401	22	AAH38458	Human neutrophil
42	16.4	82.0	401	22	AAH38459	Human neutrophil
43	16.4	82.0	401	22	AAH38460	Human neutrophil
44	16.4	82.0	401	22	AAH38461	Human neutrophil
45	16.4	82.0	401	22	AAH38462	Human neutrophil

ALIGNMENTS

AAH25706/C	standard; DNA; 836 BP.
AAH25706;	
14-AUG-2001 (first entry)	
E coli secA coding sequence.	
Antisense; microbial growth; essential gene; antimicrobial;	
Proliferation; infectious disease; secA; ds.	
Escherichia coli.	
US6228579-B1.	
08-MAY-2001.	
14-NOV-1997;	9705-0971090.
14-NOV-1997;	9705-0971090.
(UYSA-) UNIV SAN DIEGO STATE FOUND.	
Zykind JW, Forsyth RA;	
WPI; 2001-335011/35.	
Identifying microbial proliferation genes; useful for identifying	
antimicrobial agents; comprises introducing into a microorganism an	
exogenous nucleic acid having sequence identity to an endogenous	

PT microbial gene -
XX
PS Disclosure; Fig 11; 28pp; English.

CC The present invention describes a method of identifying genes essential
CC for microbial growth and proliferation, involving introducing an
CC exogenous nucleic acid into a microorganism, where the sequence is
CC similar to an endogenous microbial gene, and identifying the gene as
CC essential by comparing the organism's viability when the exogenous
CC sequence is expressed and when it is not present. This can be used to
CC identify targets for antimicrobial compounds for use in the therapy of
CC infectious diseases.

SO Sequence 836 BP; 223 A; 220 C; 198 G; 195 T; 0 other;

Query Match 100.0%; Score 20; DB 22; Length 836;
Best Local Similarity 100.0%; Pred. No. 7.4;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GATTAGCATATTAATAATCTC 20
DB 648 GATTAGCATATTAATAATCTC 629

RESULT 2

AAH25707
ID AAH25707 standard; RNA; 836 BP.

AC AAH25707;

DT 14-AUG-2001 (first entry)

DE E coli secA mRNA antisense sequence.

KW Antisense: microbial growth; essential gene; antimicrobial;
KW proliferation; infectious disease; secA; ss.

OS Escherichia coli.

XX US6228579-B1.

XX 08-MAY-2001.

XX 14-NOV-1997; 97US-0971090.

XX 14-NOV-1997; 97US-0971090.

PA (URSA-) UNITV SAN DIEGO STATE FOUND.

XX Zyskind JW, Forsyth RA.

XX WPI: 2001-335011/35.

PT Identifying microbial proliferation genes, useful for identifying
PT antimicrobial agents, comprises introducing into a microorganism an
PT exogenous nucleic acid having sequence identity to an endogenous
PT microbial gene -

PS Example 3; Fig 12; 28pp; English.

CC The present invention describes a method of identifying genes essential
CC for microbial growth and proliferation, involving introducing an
CC exogenous nucleic acid into a microorganism, where the sequence is
CC similar to an endogenous microbial gene, and identifying the gene as
CC essential by comparing the organism's viability when the exogenous
CC sequence is expressed and when it is not present. This can be used to
CC identify targets for antimicrobial compounds for use in the therapy of
CC infectious diseases.

SO Sequence 836 BP; 195 A; 198 C; 220 G; 223 U; 0 other;

Query Match 100.0%; Score 20; DB 22; Length 836;
Best Local Similarity 70.0%; Pred. No. 7.4;

Matches 14; Conservative 6; Mismatches 0; Indels 0; Gaps 0;
QY 1 GATTAGCATATTAATAATCTC 20
DB 189 GAUAGCAUUAUAAUCC 208

RESULT 3

AAH8420/C
ID AAH8420 standard; DNA; 2754 BP.

AC AAH8420;

DT 30-SEP-1999 (first entry)

DE JP1192089 Seq ID 4.

KW Tp surface antigen; secretion-related enzyme; ds.

OS Unidentified.

XX JP1192089-A.

XX 21-JUL-1999.

XX 29-DEC-1997; 97JP-0367638.

XX 29-DEC-1997; 97JP-0367638.

PA (PURE) FUJIREBIO KK.

XX WPI: 1999-461459/39.

PT Treponema pallidum-fused DNA sequence - and expression of Treponema
PT pallidum antigen by using said sequence

PS Disclosure; Page 10-12; 16pp; Japanese.

CC This invention describes a novel Treponema pallidum (Tp)-fused DNA
CC sequence in which a DNA sequence coding the surface antigen of Tp added
CC by a signal peptide is fused with a DNA sequence coding a
CC secretion-related enzyme. Also claimed is a method for expressing the
CC Tp antigen by using the above Tp-fused DNA sequence.

SO Sequence 2754 BP; 729 A; 680 C; 761 G; 584 T; 0 other;

Query Match 100.0%; Score 20; DB 20; Length 2754;
Best Local Similarity 100.0%; Pred. No. 7.3;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTAGCATATTAATAATCTC 20
DB 49 GATTAGCATATTAATAATCTC 30

RESULT 4

AAH38291/C
ID AAH38291 standard; DNA; 3811 BP.

AC AAH38291;

DT 16-JUN-1999 (first entry)

DE E. coli secA DNA.

KW Microorganism inhibitor; antisense; nuclease resistant; treatment;
KW ribonucleotide reductase; secA gene; pathological condition;
KW antimicrobial agent; crop protection; ss.

OS Escherichia coli.

XX WO9902673-A2

PD 21-JAN-1999.
XX
PF 10-JUL-1998: 98WO-CA00666.
PR
XX 10-JUL-1997: 97US-0052160.
XX
PA (GENE-) GENESENSE TECHNOLOGIES INC.
PI Dugourd D, Wright JA, Young AH;
XX WPI; 1999-120874/10.
XX
PT New oligonucleotides complementary to RR or SecA genes - useful to
XX inhibit growth of microorganisms
XX
PS Disclosure; Fig 5; 103pp; English.
XX
CC This invention describes novel antisense oligonucleotides
CC (AA38301-X38552) which are nuclease resistant, and comprises about 3-50
CC nucleotides complementary to the ribonucleotide reductase gene or the
CC secA gene of a microorganism. The antisense oligonucleotides are used to
CC treat mammalian pathological conditions mediated by microorganisms. The
CC oligonucleotides are particularly useful as antimicrobial agents in crop
CC protection. This DNA sequence contains the Escherichia coli secA gene.
XX
SQ Sequence 3811 BP; 1016 A; 942 C; 1000 G; 853 T; 0 other;

Query Match 100.0%; Score 20; DB 20; Length 3811;
Best Local Similarity 100.0%; Pred. No. 7.3;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTAGCATATATAATCTC 20
DB 830 GATTAGCATATATAATCTC 811

RESULT 5
AAC05228/C
ID AAC05228 standard; cDNA; 325 BP.
XX
AC AAC05228;
XX
DT 06-OCT-2000 (first entry)
XX
DE Human secreted protein 5' EST, SEQ ID NO: 9303.
XX
KW Human; 5' EST; expressed sequence tag; secreted protein; cDNA isolation;
KW gene therapy; chromosome mapping; ss.
XX
OS Homo sapiens.
XX
PN EP1033401-A2.
XX
PD 06-SEP-2000.
XX
PF 21-FEB-2000; 2000EP-0200610.
XX
PR 26-FEB-1999: 99US-0122487.
XX
PA (GEST) GENSET.
XX
PI Dumas Milne Edwards J, Duclert A, Giordano J;
XX WPI; 2000-500381/45.
XX
PT New nucleic acid that is a 5' expressed sequence tag (5' EST) for
PT obtaining cDNAs and genomic DNAs that correspond to 5' ESTs and for
PT diagnostic, forensic, gene therapy and chromosome mapping procedures -
XX
PS Claim 1: SEQ ID 9303; 71pp + CD-ROM; English.
XX
CC The present sequence is one of a large number of 5' ESTs derived from
CC mRNAs encoding secreted proteins. No ORF has yet been conclusively

CC identified within the present sequence. The 5' ESTs were prepared from
CC total human RNAs or polyA+ RNAs derived from 30 different tissues. EST
CC sequences usually correspond mainly to the 3' untranslated region (UTR)
CC of the mRNA because they are often obtained from oligo-dT primed cDNA
CC libraries. Such ESTs are not well suited for isolating cDNA sequences
CC derived from the 5' ends of mRNAs and even in those cases where longer
CC cDNA sequences have been obtained, the full 5' UTR is rarely included.
CC 5' ESTs are derived from mRNAs with intact 5' ends and can therefore be
CC used to obtain full length cDNAs and genomic DNAs. 5' ESTs are also used
CC in diagnostic, forensic, gene therapy and chromosome mapping procedures.
CC They are used to obtain upstream regulatory sequences and to design
CC expression and secretion vectors.
XX
SQ Sequence 325 BP; 100 A; 68 C; 68 G; 89 T; 0 other;

Query Match 92.0%; Score 18.4; DB 21; Length 325;
Best Local Similarity 95.0%; Pred. No. 37;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GATTAGCATATATAATCTC 20
DB 39 GATTAGCATATATAATCTC 20

RESULT 6
AAK61816
ID AAK61816 standard; cDNA; 645 BP.
XX
AC AAK61816;
XX
DT 06-NOV-2001 (first entry)
XX
DE Human immune/haematopoietic antigen encoding cDNA SEQ ID NO:6876.
XX
KW Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;
KW cytostatic; gene therapy; vaccine; metastasis; ss.
XX
OS Homo sapiens.
XX
PN WO200157182-A2.
XX
PD 09-AUG-2001.
XX
PF 17-JAN-2001; 2001WO-US01354.
XX
PR 31-JAN-2000; 2000US-0179065.
PR 04-FEB-2000; 2000US-0180628.
PR 24-FEB-2000; 2000US-0184664.
PR 02-MAR-2000; 2000US-0186350.
PR 16-MAR-2000; 2000US-0189874.
PR 17-MAR-2000; 2000US-0190076.
PR 18-APR-2000; 2000US-0198123.
PR 19-MAY-2000; 2000US-0205515.
PR 07-JUN-2000; 2000US-0209467.
PR 28-JUN-2000; 2000US-0214886.
PR 30-JUN-2000; 2000US-0215135.
PR 07-JUL-2000; 2000US-0216647.
PR 07-JUL-2000; 2000US-0216880.
PR 11-JUL-2000; 2000US-0217487.
PR 14-JUL-2000; 2000US-0218290.
PR 26-JUL-2000; 2000US-0220963.
PR 26-JUL-2000; 2000US-0220964.
PR 14-AUG-2000; 2000US-0224518.
PR 14-AUG-2000; 2000US-0224519.
PR 14-AUG-2000; 2000US-0225213.
PR 14-AUG-2000; 2000US-0225214.
PR 14-AUG-2000; 2000US-0225266.
PR 14-AUG-2000; 2000US-0225267.
PR 14-AUG-2000; 2000US-0225268.
PR 14-AUG-2000; 2000US-0225270.
PR 14-AUG-2000; 2000US-0225447.
PR 14-AUG-2000; 2000US-0225757.

PR	18-AUG-2000	2000US-02257559
PR	14-AUG-2000	2000US-02257558
PR	18-AUG-2000	2000US-02265779
PR	22-AUG-2000	2000US-02266681
PR	22-AUG-2000	2000US-02266681
PR	22-AUG-2000	2000US-02271882
PR	22-AUG-2000	2000US-02272079
PR	30-AUG-2000	2000US-02289824
PR	01-SEP-2000	2000US-02292987
PR	01-SEP-2000	2000US-02292987
PR	01-SEP-2000	2000US-02293444
PR	01-SEP-2000	2000US-02299345
PR	05-SEP-2000	2000US-02299509
PR	05-SEP-2000	2000US-02299513
PR	06-SEP-2000	2000US-02304337
PR	06-SEP-2000	2000US-02304338
PR	08-SEP-2000	2000US-02312482
PR	08-SEP-2000	2000US-02312482
PR	08-SEP-2000	2000US-02312444
PR	08-SEP-2000	2000US-02313113
PR	08-SEP-2000	2000US-02313114
PR	08-SEP-2000	2000US-02320800
PR	08-SEP-2000	2000US-02320801
PR	12-SEP-2000	2000US-02331968
PR	14-SEP-2000	2000US-02332997
PR	14-SEP-2000	2000US-02332998
PR	14-SEP-2000	2000US-02332999
PR	14-SEP-2000	2000US-02332999
PR	14-SEP-2000	2000US-02334201
PR	14-SEP-2000	2000US-02334201
PR	14-SEP-2000	2000US-02330663
PR	14-SEP-2000	2000US-02330665
PR	21-SEP-2000	2000US-02344223
PR	21-SEP-2000	2000US-02344223
PR	25-SEP-2000	2000US-02345997
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PR	25-SEP-2000	2000US-02358584
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PR	27-SEP-2000	2000US-02358536
PR	29-SEP-2000	2000US-02365127
PR	29-SEP-2000	2000US-02365127
PR	29-SEP-2000	2000US-02365130
PR	29-SEP-2000	2000US-02365130
PR	02-OCT-2000	2000US-02370337
PR	02-OCT-2000	2000US-02370337
PR	02-OCT-2000	2000US-02370339
PR	02-OCT-2000	2000US-02370340
PR	13-OCT-2000	2000US-02399335
PR	13-OCT-2000	2000US-02399335
PR	20-OCT-2000	2000US-02409600
PR	20-OCT-2000	2000US-02412121
PR	20-OCT-2000	2000US-02417185
PR	20-OCT-2000	2000US-02417186
PR	20-OCT-2000	2000US-02418187
PR	20-OCT-2000	2000US-02418188
PR	20-OCT-2000	2000US-02418189
PR	27-OCT-2000	2000US-02418266
PR	01-NOV-2000	2000US-02446174
PR	08-NOV-2000	2000US-02466175
PR	08-NOV-2000	2000US-02466175
PR	08-NOV-2000	2000US-02466177
PR	08-NOV-2000	2000US-02466178
PR	08-NOV-2000	2000US-02465224
PR	08-NOV-2000	2000US-02465224
PR	08-NOV-2000	2000US-02465227
PR	08-NOV-2000	2000US-02465228
PR	08-NOV-2000	2000US-02465332
PR	08-NOV-2000	2000US-02465332

PR 08-NOV-2000; 2000US-0246610.
PR 08-NOV-2000; 2000US-0246611.
PR 08-NOV-2000; 2000US-0246613.
PR 17-NOV-2000; 2000US-0249207.
PR 17-NOV-2000; 2000US-0249208.
PR 17-NOV-2000; 2000US-0249209.
PR 17-NOV-2000; 2000US-0249210.
PR 17-NOV-2000; 2000US-0249211.
PR 17-NOV-2000; 2000US-0249212.
PR 17-NOV-2000; 2000US-0249213.
PR 17-NOV-2000; 2000US-0249214.
PR 17-NOV-2000; 2000US-0249215.
PR 17-NOV-2000; 2000US-0249216.
PR 17-NOV-2000; 2000US-0249217.
PR 17-NOV-2000; 2000US-0249218.
PR 17-NOV-2000; 2000US-0249219.
PR 17-NOV-2000; 2000US-0249220.
PR 17-NOV-2000; 2000US-0249221.
PR 17-NOV-2000; 2000US-0249222.
PR 17-NOV-2000; 2000US-0249223.
PR 17-NOV-2000; 2000US-0249224.
PR 17-NOV-2000; 2000US-0249225.
PR 17-NOV-2000; 2000US-0249226.
PR 17-NOV-2000; 2000US-0249265.
PR 17-NOV-2000; 2000US-0249297.
PR 17-NOV-2000; 2000US-0249299.
PR 17-NOV-2000; 2000US-0249300.
PR 01-DEC-2000; 2000US-0250160.
PR 01-DEC-2000; 2000US-0250391.
PR 05-DEC-2000; 2000US-0251030.
PR 05-DEC-2000; 2000US-0251088.
PR 05-DEC-2000; 2000US-0256719.
PR 06-DEC-2000; 2000US-0251479.
PR 08-DEC-2000; 2000US-0251856.
PR 08-DEC-2000; 2000US-0251868.
PR 08-DEC-2000; 2000US-0251869.
PR 08-DEC-2000; 2000US-0251989.
PR 08-DEC-2000; 2000US-0251990.
PR 11-DEC-2000; 2000US-0254097.
PR 05-JAN-2001; 2001US-02595678.

(HUMA-) HUMAN GENOME SCI INC.

PA Rosen CA, Barash SC, Ruben SW;
PI WPI; 2001-483426/52.
DR P-PADB; AAM89035.

XX Nucleic acids encoding human immune/hematopoietic antigen polypeptides,
PT useful for preventing, diagnosing and/or treating cancers and
PT metastasis -

XX Claim 1; SEQ ID NO 6876; 3071pp + Sequence Listing; English.

PS AAK54951 to AAK64702 encode the human immune/hematopoietic antigen (I)
XS amino acid sequences given in AAM82170 to AAM91921. (I) have cytosolic
CC activity, and can be used in gene therapy and vaccine production. (I)
CC proteins and polynucleotides may be used in the prevention, diagnosis and
CC treatment of diseases associated with inappropriate (I) expression. For
CC example, they may be used to treat disorders associated with decreased
CC expression by rectifying mutations or deletions in a patient's genome
CC that affect the activity of (I) by expressing inactive proteins or to
CC supplement the patients own production of (I). Additionally, (I)
CC polynucleotides may be used to produce the secreted (I), by inserting
CC the nucleic acids into a host cell and culturing the cell to express the
CC protein. (I) proteins and polynucleotides may be used to prevent,
CC diagnose and treat immune/hematopoietic-related diseases, especially
CC cancers and cancer metastases of hematopoietic-derived cells. AAK64703
CC to AAK7654 represent human immune/hematopoietic antigen genomic
CC sequences from the present invention. AAK54942 to AAK54950 and AAM82169
CC represent sequences used in the exemplification of the present invention.

SQ Sequence 645 BP; 193 A; 145 C; 118 G; 186 T; 3 other;

Query Match	92.08;	Score 18.4;	DB 22;	Length 645;
Best Local Similarity	95.08;	Pred. No. 37;		
Matches 19; Conservative	0;	Mismatches 1;	Indels 0;	Gaps 0

OY 1 GATTGACATATAATATCTC 20
|||
DB 89 GAGTAGCATATAATATCTC 108

RESULT 7
AAK81074
ID AAK81074 standard; DNA: 1972 BP.

XX AAK81074;

DT 07-NOV-2001 (first entry)

DE Human Immune/Haematopoietic antigen genomic sequence SEQ ID NO:35886.

KW Human; Immune; haematopoietic; immune/haematopoietic antigen; cancer;
KW cytostatic; gene therapy; vaccine; metastasis; ds.

OS Homo sapiens.

PN WO200157182-A2.

PD 09-AUG-2001.

PF 17-JAN-2001; 2001WO-US01354.

PR 31-JAN-2000; 2000US-0179065.

PR 04-FEB-2000; 2000US-0180628.

PR 24-FEB-2000; 2000US-0184664.

PR 02-MAR-2000; 2000US-0186350.

PR 16-MAR-2000; 2000US-0189874.

PR 17-MAR-2000; 2000US-0190076.

PR 18-APR-2000; 2000US-0198123.

PR 19-MAY-2000; 2000US-0205515.

PR 07-JUN-2000; 2000US-0209467.

PR 28-JUN-2000; 2000US-0214886.

PR 30-JUN-2000; 2000US-0215135.

PR 07-JUL-2000; 2000US-0216647.

PR 07-JUL-2000; 2000US-0216880.

PR 11-JUL-2000; 2000US-0217487.

PR 11-JUL-2000; 2000US-0217496.

PR 14-JUL-2000; 2000US-0218290.

PR 26-JUL-2000; 2000US-0220963.

PR 26-JUL-2000; 2000US-0220963.

PR 14-AUG-2000; 2000US-0224518.

PR 14-AUG-2000; 2000US-0224519.

PR 14-AUG-2000; 2000US-0225213.

PR 14-AUG-2000; 2000US-0225216.

PR 14-AUG-2000; 2000US-0225266.

PR 14-AUG-2000; 2000US-0225268.

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PR 14-AUG-2000; 2000US-0225759.

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PR 01-SEP-2000; 2000US-0229345.

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PR 05-SEP-2000; 2000US-0229513.

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PR 29-SEP-2000; 2000US-0236369.

PR 29-SEP-2000; 2000US-0236370.

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PR 02-OCT-2000; 2000US-0237039.

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PR 20-OCT-2000; 2000US-0241786.

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PR 20-OCT-2000; 2000US-0241809.

PR 20-OCT-2000; 2000US-0241826.

PR 01-NOV-2000; 2000US-0244617.

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PR 08-NOV-2000; 2000US-0246478.

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PR 08-NOV-2000; 2000US-0246526.

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PR 17-NOV-2000; 2000US-0249212.

PR 17-NOV-2000; 2000US-0249213.

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PR 17-NOV-2000; 2000US-0249217.

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PR 17-NOV-2000; 2000US-0249265.

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PR	29-SEP-2000;	2000US-0236367.
PR	29-SEP-2000;	2000US-0236368.
PR	29-SEP-2000;	2000US-0236369.
PR	29-SEP-2000;	2000US-0236370.
PR	02-OCT-2000;	2000US-0236802.
PR	02-OCT-2000;	2000US-0237037.
PR	02-OCT-2000;	2000US-0237038.
PR	02-OCT-2000;	2000US-0237039.
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PR	13-OCT-2000;	2000US-0239335.
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PR	20-OCT-2000;	2000US-0240960.
PR	20-OCT-2000;	2000US-0241221.
PR	20-OCT-2000;	2000US-0241785.
PR	20-OCT-2000;	2000US-0241786.
PR	20-OCT-2000;	2000US-0241787.
PR	20-OCT-2000;	2000US-0241808.
PR	20-OCT-2000;	2000US-0241809.
PR	01-NOV-2000;	2000US-0241826.
PR	08-NOV-2000;	2000US-0244617.
PR	08-NOV-2000;	2000US-0246474.
PR	08-NOV-2000;	2000US-0246475.
PR	08-NOV-2000;	2000US-0246476.
PR	08-NOV-2000;	2000US-0246477.
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PR	08-NOV-2000;	2000US-0246524.
PR	08-NOV-2000;	2000US-0246525.
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PR	08-NOV-2000;	2000US-0246611.
PR	08-NOV-2000;	2000US-0246613.
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PR	17-NOV-2000;	2000US-0249210.
PR	17-NOV-2000;	2000US-0249211.
PR	17-NOV-2000;	2000US-0249212.
PR	17-NOV-2000;	2000US-0249213.
PR	17-NOV-2000;	2000US-0249214.
PR	17-NOV-2000;	2000US-0249215.
PR	17-NOV-2000;	2000US-0249216.
PR	17-NOV-2000;	2000US-0249217.
PR	17-NOV-2000;	2000US-0249218.
PR	17-NOV-2000;	2000US-0249244.
PR	17-NOV-2000;	2000US-0249245.
PR	17-NOV-2000;	2000US-0249264.
PR	17-NOV-2000;	2000US-0249265.
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PR	17-NOV-2000;	2000US-0249299.
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PR	05-DEC-2000;	2000US-0251030.
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PR	05-DEC-2000;	2000US-0256719.
PR	06-DEC-2000;	2000US-0251479.
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PR	08-DEC-2000;	2000US-0251868.
PR	08-DEC-2000;	2000US-0251869.
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PR	05-JAN-2001;	2001US-0259676.
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PA	(HUMA-) HUMAN GENOME SCI INC.	
XX		
PI	Rosen CA, Barash SC, Ruben SM;	

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PR 01-SEP-2000; 2000US-0229345.
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PR 05-SEP-2000; 2000US-0229513.
PR 06-SEP-2000; 2000US-0230437.
PR 06-SEP-2000; 2000US-0230438.
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PR 08-SEP-2000; 2000US-0231413.
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PR 14-SEP-2000; 2000US-0233064.
PR 14-SEP-2000; 2000US-0233065.
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PR 25-SEP-2000; 2000US-0234997.
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PR 27-SEP-2000; 2000US-0235834.
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PR 29-SEP-2000; 2000US-0236369.
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PR 01-NOV-2000; 2000US-0244617.
PR 08-NOV-2000; 2000US-0246474.
PR 08-NOV-2000; 2000US-0246475.
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PR 08-NOV-2000; 2000US-0246478.
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PR 08-NOV-2000; 2000US-0246528.
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PR 08-NOV-2000; 2000US-0246613.
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PR 17-NOV-2000; 2000US-0249211.
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PR 17-NOV-2000; 2000US-0249245.
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PR 01-DEC-2000; 2000US-0250160.
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PR 08-DEC-2000; 2000US-0251868.
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PR 08-DEC-2000; 2000US-0251990.
PR 11-DEC-2000; 2000US-0254097.
PR 05-JAN-2001; 2001US-0259678.
PR (HUMA-) HUMAN GENOME SCI INC.
PR PA
PR XX
PR XX
PR PI
PR WPI: 2001-483426/52.
PR DR
PR XX
PR PT
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PR PS
PR XX
CC AAK54951 to AAK64702 encode the human Immune/Hematopoietic antigen (I)
CC amino acid sequences given in AAK82170 to AAK91921. (I) have cytosolic
CC activity, and can be used in gene therapy and vaccine production. (I)
CC proteins and polynucleotides may be used in the prevention, diagnosis and
CC treatment of diseases associated with inappropriate (I) expression. For
CC example, they may be used to treat disorders associated with decreased
CC expression by rectifying mutations or deletions in a patient's genome
CC that affect the activity of (I) by expressing inactive proteins or to
CC supplement the patients own production of (I). Additionally, (I)
CC polynucleotides may be used to produce the secreted (I), by inserting
CC the nucleic acids into a host cell and culturing the cell to express the

protein. (i) proteins and polynucleotides may be used to prevent, diagnose and treat immune/haematopoietic-related diseases, especially cancers and cancer metastases of haematopoietic-derived cells. AAK64703 to AAK87694 represent human immune/haematopoietic antigen genomic sequences from the present invention. AAK54942 to AAK54950 and AAK82169 represent sequences used in the exemplification of the present invention.

Sequence 1976 BP; 663 A; 409 C; 350 G; 554 T; 0 other;

Query Match 92.0%; Score 18.4; DB 22; Length 1976;
Best Local Similarity 95.0%; Pred. No. 37;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GATTAGCATTAATAATCTC 20
DB 89 GAGTAGCATTAATAATCTC 108

RESULT 10
AAC59841/C
ID AAC59841 standard; DNA: 3153 BP.
AC AAC59841;
XX
XX
DT 26-JAN-2001 (first entry)
XX
XX
DE Human secreted protein encoding DNA clone vo31.1.
XX
XX
KW Secreted protein; human; autoimmune disorder; multiple sclerosis; ulcer;
KW systemic lupus erythematosus; rheumatoid arthritis; anaemia; stroke;
KW haematopoietic regulation; tissue regrowth; wound healing; haemophilia;
KW Alzheimer's disease; Parkinson's disease; Shy-Drager syndrome; cancer;
KW contraceptive; infection; growth inhibition; hyperproliferative disorder;
KW psoriasis; ds.
XX
XX
OS Homo sapiens.
XX
XX
PN WO200055375-A1.
XX
XX
PD 21-SEP-2000.
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PF 17-MAR-2000; 2000WO-US07285.
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PR 17-MAR-1999; 99US-0124808.
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PR 17-MAR-1999; 99US-0124916.
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PR 17-AUG-1999; 99US-0149639.
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PR 01-OCT-1999; 99US-0157247.
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PR 29-NOV-1999; 99US-0167824.
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PR 15-FEB-2000; 2000US-0182711.
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XX
PA (ALPH-) ALPHAGENE INC.
XX
XX
PI Valenzuela D, Yuan O, Hoffman H, Hall J, Raplejo P;
XX
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DR MPI: 2000-638211/61.
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DR P-PSDB: AAB34740.
XX
XX
XX Novel proteins and polypeptides useful for the treatment of e.g.
XX multiple sclerosis, systemic lupus erythematosus, rheumatoid arthritis,
XX cancer, Alzheimer's disease, Parkinson's disease, stroke, anemia and
XX ulcers -
XX
XX
PS Claim 116; Page 455-456; 493pp; English.
XX
XX
CC This invention relates to 59 human secreted proteins and the nucleotide
CC sequences encoding them. Sequences AAC59788-C59846 and AAB34687-B34745
CC represent the proteins and their encoding nucleotide sequences, and
CC sequences AAB34746-B34771 represent fragments of the proteins. Probes
CC for the DNA sequences are represented by sequences AAC59847-C59936. The
CC proteins exhibit neuroprotective, dermatological, immunosuppressive,
CC antiinflammatory, antianemic, nootropic, antiparkinsonian,
CC cerebroprotective, haemostatic, vulnery, cytostatic, antipsoriatic,
CC antibacterial, virucide, and fungicide activity. The proteins and

nucleotide sequences are useful as nutritional sources or supplements
CC and in research. The proteins are useful for treating immune deficiency
CC and disorders, which may be genetic or resulting from infections,
CC autoimmune disorders such as multiple sclerosis, systemic lupus
CC erythematosus, rheumatoid arthritis, and for treating myeloid or lymphoid
CC cell deficiencies such as anaemias by regulating haematopoiesis. The
CC proteins are also useful in compositions for bone, cartilage, tendon,
CC ligament and/or nerve tissue growth or regeneration, for wound healing,
CC tissue repair and replacement and in the treatment of wounds, incisions
CC and ulcers. Other uses include in the treatment of central and
CC peripheral nervous system and neuropathies such as Alzheimer's and
CC Parkinson's diseases and Shy-Drager syndrome, and mechanical and
CC traumatic disorders, such as spinal cord disorders, head trauma and
CC stroke. The proteins may also be used as a contraceptive, and for
CC treating coagulation disorders such as haemophilias. The protein and
CC nucleotide sequences with cadherin activity are useful for treating
CC cancer. Other uses for the protein include for inhibiting the growth,
CC infection or function of, or killing, infectious agents such as bacteria,
CC virus, fungi and other parasites, for effecting bodily characteristics
CC such as height, weight, hair colour, effecting biorhythms or cardiac
CC cycles or rhythms, effecting metabolism, catabolism, anabolism,
CC processing, utilization, storage or elimination of dietary fat, lipid,
CC protein, carbohydrate, vitamins, minerals, cofactors, effecting
CC behavioural characteristics, providing analgesic effects and for treating
CC hyperproliferative disorders such as psoriasis.

Sequence 3153 BP; 938 A; 709 C; 647 G; 859 T; 0 other;

Query Match 87.0%; Score 17.4; DB 21; Length 3153;
Best Local Similarity 94.7%; Pred. No. 1e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 ATTAGCATTAATAATCTC 20
DB 1944 ATTGGCATTAATAATCTC 1926

RESULT 11
AAK95332/C
ID AAK95332 standard; DNA: 401 BP.
XX
XX
AC AAK95332;
XX
XX
DT 17-DEC-2001 (first entry)
XX
XX
DE Human neuregulin gene single nucleotide polymorphism SNP8NRG6787.
XX
XX
KW Human: neuregulin-1 associated gene 1; NRG1AG1; Schizophrenia gene;
KW gene therapy; single nucleotide polymorphism; SNP; ds.
XX
XX
OS Homo sapiens.
XX
XX
PN WO200164876-A2.
XX
XX
PD 07-SEP-2001.
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PF 28-FEB-2001; 2001WO-US06376.
XX
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PR 28-FEB-2000; 2000US-0515715.
XX
XX
PA (DECO-) DECODE GENETICS EHF.
XX
XX
PI Stefansson H, Steinthorsdottir V, Gulcher JR;
XX
XX
DR MPI: 2001-550179/61.
XX
XX
PT Neuregulin-1 associated gene 1 nucleic acids and fragments, useful for
XX preventing diagnosing and treating schizophrenia -
XX
XX
PS Disclosure; Page 510; 750pp; English.
XX
XX
CC This sequence represents a single nucleotide polymorphism (SNP) of the
CC human neuregulin-1 associated gene 1 (NRG1AG1) of the invention. The

CC NR1A1 gene is also referred to as the human Schizophrenia gene. The
CC invention also relates to fragments or variants of the gene and the
CC NR1A1 polypeptides they encode. The NR1A1 nucleic acids and
CC polypeptides may be used in the prevention, diagnosis and treatment of
CC diseases associated with inappropriate NR1A1 expression. For example,
CC they may be used to treat disorders associated with decreased expression
CC by rectifying mutations or deletions in a patient's genome that affect
CC the activity of NR1A1 by expressing inactive proteins or to supplement
CC the patients own production of NR1A1. Additionally, the gene may be
CC used to produce NR1A1 polypeptides, by inserting the nucleic acids into
CC a host cell and culturing the cell to express the protein. The gene may
CC also be used as DNA probes and primers in diagnostic assays to detect and
CC quantitate the presence of similar nucleic acids in samples, and
CC therefore which patients may be in need of restorative therapy. The
CC NR1A1 polypeptides may also be used as antigens in the production of
CC antibodies against NR1A1 and in assays to identify modulators of
CC NR1A1 expression and activity. Anti-NR1A1 antibodies and antagonists
CC may also be used to down regulate expression and activity. Anti-NR1A1
CC antibodies may also be used as diagnostic agents for detecting the
CC presence of NR1A1 polypeptides in samples. NR1A1 is associated with
CC schizophrenia which may be prevented, diagnosed and/or treated by the
CC above methods.

XX Sequence 401 BP; 119 A; 96 C; 84 G; 98 T; 4 other;

Query Match 82.0%; Score 16.4; DB 22; Length 401;

Best Local Similarity 94.4%; Pred. No. 2.8e+02;

Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GATTAGCATATTAATC 18

DB 358 GATTGCGATTAATAATC 341

RESULT 12

AAK95333/c

ID AAK95333 standard; DNA; 401 BP.

XX AAK95333;

DT 17-DEC-2001 (first entry)

DE Human neuregulin gene single nucleotide polymorphism SNP8NRG6844.

KW Human; neuregulin-1 associated gene 1; NR1A1; Schizophrenia gene;

KM gene therapy; single nucleotide polymorphism; SNP; ds.

OS Homo sapiens.

PN WO200164876-A2.

PD 07-SEP-2001.

PF 28-FEB-2001; 2001WO-US06376.

PR 28-FEB-2000; 2000US-0515715.

PA (DECO-) DECODE GENETICS EHF.

PI Stefansson H, Steinthorsdottir V, Gulcher JR;

DR WPI; 2001-550179/61.

PT Neuregulin-1 associated gene 1 nucleic acids and fragments, useful for

PT preventing diagnosing and treating schizophrenia -

XX Disclosure; Page 510; 750pp; English.

XX This sequence represents a single nucleotide polymorphism (SNP) of the
CC human neuregulin-1 associated gene 1 (NR1A1) of the invention. The
CC NR1A1 gene is also referred to as the human Schizophrenia gene. The
CC invention also relates to fragments or variants of the gene and the
CC NR1A1 polypeptides they encode. The NR1A1 nucleic acids and

CC polypeptides may be used in the prevention, diagnosis and treatment of
CC diseases associated with inappropriate NR1A1 expression. For example,
CC they may be used to treat disorders associated with decreased expression
CC by rectifying mutations or deletions in a patient's genome that affect
CC the activity of NR1A1 by expressing inactive proteins or to supplement
CC the patients own production of NR1A1. Additionally, the gene may be
CC used to produce NR1A1 polypeptides, by inserting the nucleic acids into
CC a host cell and culturing the cell to express the protein. The gene may
CC also be used as DNA probes and primers in diagnostic assays to detect and
CC quantitate the presence of similar nucleic acids in samples, and
CC therefore which patients may be in need of restorative therapy. The
CC NR1A1 polypeptides may also be used as antigens in the production of
CC antibodies against NR1A1 and in assays to identify modulators of
CC NR1A1 expression and activity. Anti-NR1A1 antibodies and antagonists
CC may also be used to down regulate expression and activity. Anti-NR1A1
CC antibodies may also be used as diagnostic agents for detecting the
CC presence of NR1A1 polypeptides in samples. NR1A1 is associated with
CC schizophrenia which may be prevented, diagnosed and/or treated by the
CC above methods.

XX Sequence 401 BP; 114 A; 89 C; 92 G; 103 T; 3 other;

Query Match 82.0%; Score 16.4; DB 22; Length 401;

Best Local Similarity 94.4%; Pred. No. 2.8e+02;

Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GATTAGCATATTAATC 18

DB 301 GATTGCGATTAATAATC 284

RESULT 13

AAK96579/c

ID AAK96579 standard; DNA; 401 BP.

XX AAK96579;

DT 17-DEC-2001 (first entry)

DE Human neuregulin gene insertion/deletion DNP8NRG1.

KW Human; neuregulin-1 associated gene 1; NR1A1; Schizophrenia gene;

KM gene therapy; insertion; deletion; ds.

OS Homo sapiens.

PN WO200164876-A2.

PD 07-SEP-2001.

PF 28-FEB-2001; 2001WO-US06376.

PR 28-FEB-2000; 2000US-0515715.

PA (DECO-) DECODE GENETICS EHF.

PI Stefansson H, Steinthorsdottir V, Gulcher JR;

DR WPI; 2001-550179/61.

PT Neuregulin-1 associated gene 1 nucleic acids and fragments, useful for

PT preventing diagnosing and treating schizophrenia -

XX Disclosure; Page 729; 750pp; English.

XX This sequence represents an insertion/deletion variant of the human
CC neuregulin-1 associated gene 1 (NR1A1) of the invention. The NR1A1
CC gene is also referred to as the human Schizophrenia gene. The invention
CC also relates to fragments or variants of the gene and the NR1A1
CC polypeptides they encode. The NR1A1 nucleic acids and polypeptides may
CC be used in the prevention, diagnosis and treatment of diseases associated
CC with inappropriate NR1A1 expression. For example, they may be used to
CC treat disorders associated with decreased expression by rectifying

CC mutations or deletions in a patient's genome that affect the activity of
CC NRGIAG1 by expressing inactive proteins or to supplement the patients own
CC production of NRGIAG1. Additionally, the gene may be used to produce
CC NRGIAG1 polypeptides, by inserting the nucleic acids into a host cell
CC and culturing the cell to express the protein. The gene may also be used
CC as DNA probes and primers in diagnostic assays to detect and quantitate
CC the presence of similar nucleic acids in samples, and therefore which
CC patients may be in need of restorative therapy. The NRGIAG1 polypeptides
CC may also be used as antigens in the production of antibodies against
CC NRGIAG1 and in assays to identify modulators of NRGIAG1 expression and
CC activity. Anti-NRGIAG1 antibodies and antagonists may also be used to
CC down regulate expression and activity. Anti-NRGIAG1 antibodies may
CC also be used as diagnostic agents for detecting the presence of NRGIAG1
CC polypeptides in samples. NRGIAG1 is associated with schizophrenia which
CC may be prevented, diagnosed and/or treated by the above methods.

SO Sequence 401 BP; 124 A; 78 C; 80 G; 117 T; 2 other;

Query Match 82.0%; Score 16.4; DB 22; Length 401;
Best Local Similarity 94.4%; Pred. No. 2.8e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GATTACCATATAAATC 18
|||||
DB 186 GATTTCATATAAATC 169

RESULT 14
AAK96581/C
ID AAK96581 standard; DNA; 401 BP.

XX AAK96581;

DT 17-DEC-2001 (first entry)

DE Human neuregulin gene insertion/deletion DNP8NRG3.

KW Human; neuregulin-1 associated gene 1; NRGIAG1; Schizophrenia gene;
KW gene therapy; Insertion; deletion; ds.

OS Homo sapiens.

PN WO200164876-A2.

PD 07-SEP-2001.

PF 28-FEB-2001; 2001WO-US06376.

PR 28-FEB-2000; 2000US-0515715.

PA (DECO-) DECODE GENETICS EHF.

PI Stefansson H, Steinthorsdottir V, Gulcher JR;

DR WPI; 2001-550179/61.

PT Neuregulin-1 associated gene 1 nucleic acids and fragments, useful for
PT preventing diagnosing and treating schizophrenia

PS Disclosure; Page 729; 750pp; English.

CC This sequence represents an insertion/deletion variant of the human
CC neuregulin-1 associated gene 1 (NRGIAG1) of the invention. The NRGIAG1
CC gene is also referred to as the human Schizophrenia gene. The invention
CC also relates to fragments or variants of the gene and the NRGIAG1
CC polypeptides they encode. The NRGIAG1 nucleic acids and polypeptides may
CC be used in the prevention, diagnosis and treatment of diseases associated
CC with inappropriate NRGIAG1 expression. For example, they may be used to
CC treat disorders associated with decreased expression by rectifying
CC mutations or deletions in a patient's genome that affect the activity of
CC NRGIAG1 by expressing inactive proteins or to supplement the patients own
CC production of NRGIAG1. Additionally, the gene may be used to produce
CC NRGIAG1 polypeptides, by inserting the nucleic acids into a host cell

CC and culturing the cell to express the protein. The gene may also be used
CC as DNA probes and primers in diagnostic assays to detect and quantitate
CC the presence of similar nucleic acids in samples, and therefore which
CC patients may be in need of restorative therapy. The NRGIAG1 polypeptides
CC may also be used as antigens in the production of antibodies against
CC NRGIAG1 and in assays to identify modulators of NRGIAG1 expression and
CC activity. Anti-NRGIAG1 antibodies and antagonists may also be used to
CC down regulate expression and activity. Anti-NRGIAG1 antibodies may
CC also be used as diagnostic agents for detecting the presence of NRGIAG1
CC polypeptides in samples. NRGIAG1 is associated with schizophrenia which
CC may be prevented, diagnosed and/or treated by the above methods.

SO Sequence 401 BP; 119 A; 74 C; 85 G; 123 T; 0 other;

Query Match 82.0%; Score 16.4; DB 22; Length 401;
Best Local Similarity 94.4%; Pred. No. 2.8e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GATTACCATATAAATC 18
|||||
DB 90 GATTTCATATAAATC 73

RESULT 15
AAK96825/C
ID AAK96825 standard; DNA; 401 BP.

XX AAK96825;

DT 17-DEC-2001 (first entry)

DE Human neuregulin gene single nucleotide polymorphism SNP8NRG6787.

KW Human; neuregulin 1 gene; schizophrenia; gene therapy; SNP;

KW single nucleotide polymorphism; ds.

OS Homo sapiens.

PN WO200164877-A2.

PD 07-SEP-2001.

PF 28-FEB-2001; 2001WO-US06377.

PR 28-FEB-2000; 2000US-0515716.

PA (DECO-) DECODE GENETICS EHF.

PI Stefansson H, Steinthorsdottir V, Gulcher JR;

DR WPI; 2001-514841/56.

PT Neuregulin 1 nucleic acids and proteins useful for diagnosing
PT preventing and treating schizophrenia

PS Disclosure; Page 95; 756pp; English.

CC This sequence represents a single nucleotide polymorphism (SNP)
CC from the human neuregulin 1 gene of the invention.
CC The invention also relates to fragments or variants of the neuregulin 1
CC gene. The gene and its proteins may be used in the prevention, diagnosis
CC and treatment of diseases associated with inappropriate neuregulin 1
CC expression, such as schizophrenia. For example they may be used to treat
CC disorders associated with decreased neuregulin 1 expression by rectifying
CC mutations or deletions in a patient's genome that affect the activity of
CC neuregulin 1 by expressing inactive proteins or to supplement the
CC patients own production of polypeptides. Additionally, the gene may be
CC used to produce the neuregulin 1 protein, by inserting the nucleic acids
CC into a host cell and culturing the cell to express the protein. The gene
CC and its complementary sequences may also be used as DNA probes in
CC diagnostic assays to detect and quantitate the presence of similar
CC nucleic acids in samples, and therefore which patients may be in need of
CC restorative therapy. The protein may also be used as antigens in the

CC production of antibodies against neuregulin 1 and in assays to identify
 CC modulators of neuregulin 1 expression and activity. The antibodies and
 CC antagonists may also be used to down regulate expression and activity.
 CC The antibodies may also be used as diagnostic agents for detecting the
 CC presence of neuregulin 1 in samples.

XX
 SQ Sequence 401 BP; 119 A; 96 C; 84 G; 98 T; 4 other;

Query Match 82.0%; Score 16.4; DB 22; Length 401;
 Best Local Similarity 94.4%; Pred. No. 2.8e+02;
 Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GATTAGCATATATAATC 18
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 DB 358 GATTGACATATAATAATC 341

Search completed: October 6, 2002, 15:21:06
 Job time : 290 secs

GenCore version 5.1.3
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OM nucleic - nucleic search, using SW model

Run on: October 6, 2002, 13:13:14 ; Search time 2261 Seconds
(without alignments)
185.109 Million cell updates/sec

Title: US-09-754-468-47
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Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 1797656 seqs, 10463268293 residues
Total number of hits satisfying chosen parameters: 3595312

Minimum DB seq length: 0
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Post-processing: Minimum Match 08
Maximum Match 1008
Listing first 45 summaries

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2: gb_hng:*
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10: gb_ro:*
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12: gb_sy:*
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17: em_hum:*
18: em_in:*
19: em_mu:*
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31: em_hng_inv:*
32: em_hng_other:*
33: em_hngc_inv:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No. Score Query Match Length DB ID Description

1	C	20	100.0	20	6	AX191765	AX191765 Sequence
2	C	20	100.0	836	6	ARI49152	ARI49152 Sequence
3	C	20	100.0	836	6	ARI49153	ARI49153 Sequence
4	C	20	100.0	2048	1	ECOEENVA	M19211 E.coli cell
5	C	20	100.0	2754	6	E27365	E27365 Treponema P
6	C	20	100.0	3811	1	ECOSECA	M20791 Escherichia
7	C	20	100.0	12434	1	AE000119	AE000119 Escherich
8	C	20	100.0	12518	1	AE005186	AE005186 Escherich
9	C	20	100.0	28277	1	EC2MIN	X55034 E. coli 2 m
10	C	20	100.0	28277	6	AX191720	AX191720 Sequence
11	C	20	100.0	111401	1	EC0110K	D10483 E.coli K12
12	C	19	95.0	155976	2	AC055113	AC055113 Homo sapi
13	C	19	95.0	179212	9	AC084877	AC084877 Homo sapi
14	C	14	92.0	22286	1	AE008700	AE008700 Salmonell
15	C	15	92.0	127603	9	AC036102	AC036102 Homo sapi
16	C	16	92.0	138846	2	AC068478	AC068478 Homo sapi
17	C	18	92.0	156978	2	AC023379	AC023379 Homo sapi
18	C	18	92.0	157385	9	AC006022	AC006022 Homo sapi
19	C	18	92.0	165102	2	AC074279	AC074279 Homo sapi
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24	C	24	92.0	188049	2	AC025370	AC025370 Homo sapi
25	C	25	92.0	193588	2	AC092433	AC092433 Homo sapi
26	C	26	92.0	194782	9	AC079271	AC079271 Homo sapi
27	C	27	92.0	200475	2	AL158047	AL158047 Human DNA
28	C	28	92.0	201629	2	AC013452	AC013452 Homo sapi
29	C	29	92.0	204790	9	AL627265	AL627265 Salmonell
30	C	30	92.0	251050	1	AF363578	AF363578 Homo sapi
31	C	31	92.0	347253	2	AC094868	AC094868 Rattus no
32	C	32	90.0	51869	2	AL359976	AL359976 Homo sapi
33	C	33	90.0	175456	2	AC074364	AC074364 Homo sapi
34	C	34	90.0	195290	2	AC038729	AC038729 Giardia 1
35	C	35	94.2	942	33	AC061176	AC061176 Giardia 1
36	C	36	87.0	63769	2	AC079334	AC079334 Homo sapi
37	C	37	87.0	73610	2	AL359735	AL359735 Human DNA
38	C	38	87.0	102990	2	AP000801	AP000801 Homo sapi
39	C	39	87.0	109568	2	AP000840	AP000840 Homo sapi
40	C	40	87.0	126755	9	HSB18C9	AL049709 Human DNA
41	C	41	87.0	138969	9	AC060771	AC060771 Homo sapi
42	C	42	87.0	142278	2	AC005177	AC005177 Homo sapi
43	C	43	87.0	150406	2	AC098684	AC098684 Mus muscu
44	C	44	87.0	161742	2	AC036204	AC036204 Homo sapi
45	C	45	87.0	161742	2	AC036204	AC036204 Homo sapi

ALIGNMENTS

RESULT 1
AX191765
LOCUS AX191765 20 bp DNA linear PAT 15-AUG-2001
DEFINITION Sequence 47 from Patent WO0149775.
ACCESSION AX191765
VERSION AX191765.1 GI:15209934
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1 (bases 1 to 20)
AUTHORS Iversen, P.L.
TITLE Antisense antibacterial cell division composition and method
JOURNAL Patent: WO 0149775-A 47 12-JUL-2001;
Avi Biopharma, Inc. (US)
FEATURES
source location/Qualifiers
1..20
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/note="Oligonucleotide"

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Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 GATTAGCATATAAATCTC 20
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Db 1 GATTAGCATATAAATCTC 20

RESULT 2
LOCUS ARI49152 836 bp DNA linear PAT 08-AUG-2001
DEFINITION Sequence 6 from patent US 6228579.
ACCESSION ARI49152
VERSION ARI49152.1 GI:15113743
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 836)
AUTHORS Zyskind,J.W. and Forsyth,R.Allyn.
TITLE Method for identifying microbial proliferation genes
JOURNAL Patent: US 6228579-A 6 08-MAY-2001;
FEATURES
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BASE COUNT 223 a 220 c 198 g 195 t
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Query Match          100.0%; Score 20; DB 6; Length 836;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 GATTAGCATATAAATCTC 20
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Db 648 GATTAGCATATAAATCTC 629

RESULT 3
LOCUS ARI49153 836 bp DNA linear PAT 08-AUG-2001
DEFINITION Sequence 7 from patent US 6228579.
ACCESSION ARI49153
VERSION ARI49153.1 GI:15113744
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 836)
AUTHORS Zyskind,J.W. and Forsyth,R.Allyn.
TITLE Method for identifying microbial proliferation genes
JOURNAL Patent: US 6228579-A 7 08-MAY-2001;
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BASE COUNT 195 a 198 c 220 g 223 t
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Query Match          100.0%; Score 20; DB 6; Length 836;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 GATTAGCATATAAATCTC 20
    |||
Db 189 GATTAGCATATAAATCTC 208

RESULT 4
LOCUS ECOENVA/c 2048 bp ss-DNA linear BCT 20-DEC-1995
DEFINITION E.coli cell permeability-cell separation protein (enva) gene,
complete cds., fts2 gene, 3' end, and secA gene, 5' end.
ACCESSION M19211

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VERSION M19211.1 GI:145846
KEYWORDS cell permeability-cell separation protein; enva gene; fts2 gene;
secA gene.
SOURCE Escherichia coli (strain K-12) (clone: PACYC184.) DNA.
ORGANISM Escherichia coli
REFERENCE Beall,B. and Lutkenhaus,J.
AUTHORS Sequence analysis, transcriptional organization, and insertional
TITLE mutagenesis of the enva gene of Escherichia coli
JOURNAL J. Bacteriol. 169 (12), 5408-5415 (1987)
MEDLINE 88058745
COMMENT Draft entry and computer readable of sequence [1] kindly provided
by J. Lutkenhaus (19-FEB-1988). (put. see kay for more info.).
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Best Local Similarity 100.0%; Pred. No. 88;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 GATTAGCATATAAATCTC 20

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Db 1855 GATTACCATTAATAATCTC 1836
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RESULT 5 27365 2754 bp DNA linear PAT 07-FEB-2001
LOCUS E27365/c Treponema pallidum-fused DNA sequence and method for expressing T.
DEFINITION pallidum antigen with the use of the said sequence.
ACCESSION E27365
VERSION E27365.1 GI:13018177
KEYWORDS JP 1999192089-A/4.
SOURCE unclassified.
ORGANISM unclassified.
REFERENCE 1 (bases 1 to 2754)
AUTHORS Katsuya, F., Y.H.H. and Ito.
TITLE Treponema pallidum-fused DNA sequence and method for expressing T.
JOURNAL pallidum antigen with the use of the said sequence
PATENT: JP 1999192089-A 4 21-JUL-1999.
FUJIREBIO INC
COMMENT OS Unidentified
PN JP 1999192089-A/4
PD 21-JUL-1999
PE 29-DEC-1997 JP 1997367638
PR
PI KATSUYA FUJIMURA, YASUHIRO HARA, SATOSHI ITO
PC C12N15/09/C07K14/20,C07K19/00,C12N1/21,C12P21/02,G01N33/53,
PC G01N33/577,
PC C12N15/09,C12R1:01),(C12N1/21,C12R1:19),(C12P21/02,C12R1:19),
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PC C12N15/00,C12R1:01)
CC Strandedness: Double;
CC Topology: Linear;
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FT source /organism='Unidentified'.
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1. 2754 /organism='unidentified'
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Best Local Similarity 100.0%; Pred. No. 83;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 1 GATTACCATTAATAATCTC 20
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Db 49 GATTACCATTAATAATCTC 30
RESULT 6
LOCUS ECOSECA 3811 bp DNA linear BCT 26-APR-1993
DEFINITION Escherichia coli SecA protein gene, complete cds; mult 5' end.
ACCESSION M20791
VERSION M20791.1 GI:147792
KEYWORDS secA protein.
SOURCE E.coli (strain MC4100) DNA.
ORGANISM Escherichia coli
Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;
Escherichia.
REFERENCE 1 (bases 1 to 3811) Grodberg, I. and Oliver, D.
AUTHORS Schmidt, M., Rollo, E., and Grodberg, I. and Oliver, D.
TITLE Nucleotide sequence of secA gene and secA(ts) mutations preventing
protein export in Escherichia coli
JOURNAL J. Bacteriol. 170, 3404-3414 (1988)
MEDLINE 88298644
COMMENT Draft entry and computer readable sequence [1] kindly submitted by
M. Schmidt 28-SEP-1988
The mult gene was identified in Mcl. Gen. Genet. 206, 9-16 (1987)

FEATURES accession number X04831.
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ORIGIN 3 bp upstream of PvuII site.
Query Match 100.0%; Score 20; DB 1; Length 3811;
Best Local Similarity 100.0%; Pred. No. 78;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 1 GATTACCATTAATAATCTC 20
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Db 830 GATTACCATTAATAATCTC 811
RESULT 7
LOCUS AE000119/c 12434 bp DNA linear BCT 01-DEC-2000
DEFINITION Escherichia coli K12 MG1655 section 9 of 400 of the complete
genome.
ACCESSION AE000119 U000096
VERSION AE000119.1 GI:1786283
KEYWORDS
SOURCE Escherichia coli K12.
ORGANISM Escherichia coli K12
Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;
Escherichia.

[illegible]

Query Match	Best Local Similarity	100.0%	Score 20;	DB 1;	Length 12434;		
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DEFINITION							
ACCESSION							
VERSION							
SOURCE							
ORGANISM							
REFERENCE							
AUTHORS							

TITLE	Genome sequence of enterohaemorrhagic Escherichia coli O157:H7
JOURNAL	Nature 409 (819), 529-533 (2001)
MEDLINE	21074935
PUBMED	11206551
REFERENCE	2 (bases 1 to 12518)
AUTHORS	Perna,N.T., Plunkett,G. III, Burland,V., Mau,B., Glasner,J.D., Rose,D.J., Mayhew,G.F., Evans,P.S., Gregor,J., Kirkpatrick,H.A., Postal,G., Hackett,J., Klink,S., Boutin,A., Shao,Y., Miller,L., Grobeck,E.J., Davis,N.W., Lim,A., Dimantanta,E., Potamouzis,K., Apodaca,T., Anantharaman,T.S., Ian,J., Yen,G., Schwartz,D.C., Welch,R.A. and Blattner,F.R.
TITLE	Direct Submission
JOURNAL	Submitted (22-OCT-2000) Laboratory of Genetics, University of Wisconsin,
FEATURES	Location/Qualifiers 445 Henry Mall, Madison, WI 53706, USA
SOURCE	1..12518 organism="Escherichia coli O157:H7 EDL933" strain="EDL933" serotype="O157:H7" db_xref="taxon:15864" note="enterohemorrhagic"
gene	38..1513 /gene="murC" /note="Z0101" 38..1513 /function="enzyme; Cell envelop: Murein sacculus, peptidoglycan" /note="Residues 1 to 491 of 491 are 99.79 pct identical to residues 1 to 491 of 491 from Escherichia coli K-12 Strain MG1655: B0091" /codon_start=1 /transl_table=11 /product="L-alanine adding enzyme, UDP-N-acetyl-muramate:alanine ligase" /db_xref="GI:12512797" /translation="MNTQOLAKRSIVPEMKRRVRIHFVGIGAGMGIAEVLANEQY QISSDIAPNPVTQQLMLGATIFYFNRPENVPDASVVSSAISADNPELVAAHEAR IPVRRAEMELAEFRPHGTAIACHTGGTKTTAVASSIYAEGDPFENGWKAAGA VHARGRGLTIAEDSDASFILLOPMVAIIYNTEADHDOTGDGEDEIKOPINFL HNLPFYRAYWCDDPYIRIELPRVYGKQTITTFGSSEADVREDYVOIGRGHTTLR QDEPKRVLTSLAMPGRNALNAAVAATEBGIDDEAIDLRLAESFGTGREFDLGEFE PLEVNGVSGTAGPVLDYVDGHPTVEDATTAAARAQGWPDKNLMLEPQHRTFRDLVD DFANVLTPQDTLLMLEVYPAGEAPIPGADSRSCLORTRGRCIDPIVPPAQVAEWML APVLTGNDILTVOGAGNIKGRIARSIAIEIKLKPPREEQHD"
CDS	1506..2426 /gene="ddIB" /note="Z0102" 1506..2426 /function="enzyme; Cell envelop: Murein sacculus, peptidoglycan" /note="Residues 1 to 306 of 306 are 99.34 pct identical to residues 1 to 306 of 306 from Escherichia coli K-12 Strain MG1655: B0092" /codon_start=1 /transl_table=11 /product="D-alanine-D-alanine ligase B, affects cell division" /protein_id="AAG54396.1" /db_xref="GI:12512798" /translation="MTDKIALVIGGSTSAEREVSLSNGCAAVLAGREGIDAYPVDPKE VDVTAQSKMGFOKFALHGSGGEDGLGMLTELGMPLPYGSGYMALSMDXLRSLKL LWGCGAPPAVPAVALTYVEEFKGISDQLAETSLAGLPVLYKPRESSVGMKSYAB NALQDALRLFQHDEEVLIERWLSPGFVAAILGEELPSVRIQSTGYDYEKYLUS DETGYFCPAGLEASQEANLQALVLAKTATLGCCKMGRIDWLSDGOFYLEANTSPG MTGSIVLMAAPAQAQGMSQSOLVVIIEILD"
gene	2428..3258 /gene="ftsQ" /note="Z0103" 2428..3258 /gene="ftsQ" /note="Z0103" 2428..3258 /gene="ftsQ"
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/function="phenotype; Cell division"
/note="Residues 1 to 276 of 276 are 99.63 pct identical to
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MG1655: B0093"
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/transl_table="MSQAALNTRNSEESSRRNGTBLAGLFLITLVITYLVSGV
VLGMEADQRLPLSLVLTGERHYTRNDIRQSLILALDEPTFTQDVIITQIQEOR
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/note="20104"
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Strain MG1655: B0094"
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process, complexes with FtsZ, associated with junctions of
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ITCHMDAKNITVAVERGCLVNDLIIFAGLASSSVLTEDRELGCYVDIGGTMDI
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/note="20105"
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/gene="ftsZ"
/function="enzyme; Cell division"
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to residues 1 to 383 of 383 from Escherichia coli K-12
Strain MG1655: B0095"
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/product="cell division: forms circumferential ring;
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ITIPNDKLLVYRGISILIDAFGANVDKLGAVOCIAILIRPGAMNDFDRVMS
EMGVAMGSGVASGEDRAEAEKMAISSPLELIDSGARGLVNTITGFDRLDEFE
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Strain MG1655: B0096"
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6903.7490
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/gene="yacA"
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MG1655: B0097"
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/protein_id="AAG54401.1"
/db_xref="GI:12512803"
/transl_table="MLWTSGFNDKICALNTEFEYDRDGNVSGIITRMPOGKRYFWPH
LLGWNVASLGLPALSNAPAPAKATTRNHEPSAKVFCOLALLLEANTRRPNYS
VDYHOHAIIRYIRHLSPFAMAPOITPAVEESIPIQAOHALLDITSLALTDGPTSEK
GYRIDAHFTQAKSTFVWISQAGIRAGFORLS"
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7552.10257
/gene="seca"
/function="transport; Transport of large molecules:
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/note="Residues 1 to 901 of 901 are 99.88 pct identical to
residues 1 to 901 of 901 from Escherichia coli K-12 Strain
MG1655: B0098"
/codon_start=1
/transl_table=11
/product="transport; Transport of large molecules:
protein, peptide secretion"
/note="Residues 1 to 901 of 901 are 99.88 pct identical to
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MG1655: B0098"
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Query Match 100.0%; Score 20; DB 1; Length 12518;
Best Local Similarity 100.0%; Pred. No. 62;
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Oy 1 GATTACGATTAATAAATCTC 20
Db 7560 GATTACGATTAATAAATCTC 7541

RESULT 9
EC2MIN 28277 bp DNA linear BCT 05-JUL-1999
LOCUS EC2MIN/c
DEFINITION E. coli 2 minute region.
ACCESSION X55034 M10429
VERSION X55034.1 GI:40841
KEYWORDS
ddl gene; envA gene; ftsA gene; ftsQ gene; ftsW gene; ftsZ gene;
ilvH gene; ilvI gene; leuA gene; leuO gene; murJ gene; murC gene;
murD gene; murE gene; murF gene; murG gene; murH gene; murI gene;
orfC; orfX; pppB gene; seca gene; shl gene.
SOURCE Escherichia coli.
ORGANISM Escherichia coli.
Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;
Escherichia.
REFERENCE 1 (bases 1 to 306)
AUTHORS Wessler,S.R. and Calvo,J.M.
TITLE Control of leu operon expression in Escherichia coli by a
transcription attenuation mechanism
J. Mol. Biol. 149, 579-579 (1981)
JOURNAL MEDLINE 82078077
2 (bases 2101 to 4431)
REFERENCE Squares,C.H., Defelice,M., Devereux,J. and Calvo,J.M.
AUTHORS Molecular structure of ilvH and its evolutionary relationship to
TITLE ilvE in Escherichia coli
JOURNAL Nucleic Acids Res. 11, 5299-5313 (1983)
MEDLINE 83272971

```

- REFERENCE 3 (bases 7316 to 10074)
AUTHORS Nakamura,M., Maruyama,I.N., Some,M., Kato,J.I., Suzuki,H. and Hirota,Y.
TITLE On the process of cellular division in *Escherichia coli*: Nucleotide sequence of the gene for penicillin-binding protein 3
JOURNAL Mol. Gen. Genet. 191, 1-9 (1983)
MEDLINE 83296957
REFERENCE 4 (bases 19464 to 21952)
AUTHORS Robinson,A.C., Keenan,D.J., Hatfull,G.F., Sullivan,N.F., Spiegelberg,R. and Donachie,W.D.
TITLE DNA sequence and transcriptional organization of essential cell division genes *ftsQ* and *ftsA* of *Escherichia coli*: Evidence for overlapping transcriptional units
J. Bacteriol. 160, 546-555 (1984)
MEDLINE 85054557
REFERENCE 5 (bases 21464 to 23333)
AUTHORS Yi,Q.M. and Lutkenhaus,J.
TITLE The nucleotide sequence of the essential cell-division gene *ftsZ* of *Escherichia coli*
J. Bacteriol. 166, 1113-1117 (1986)
MEDLINE 86083166
REFERENCE 6 (bases 1799 to 2187)
AUTHORS Hauglin,G.W., Squitres,C.H., Defelice,M., Largo,C.T. and Calvo,J.M.
TITLE Unusual organization of the *llyV* promoter of *Escherichia coli*
J. Bacteriol. 163, 186-198 (1985)
MEDLINE 85234358
REFERENCE 7 (bases 268 to 1130)
AUTHORS Hauglin,G.W., Wessler,S.R., Gemmill,R.M. and Calvo,J.M.
TITLE High A + T content conserved in DNA sequences upstream of *leuABCD* in *Escherichia coli* and *Salmonella typhimurium*
J. Bacteriol. 166, 1113-1117 (1986)
MEDLINE 86223773
REFERENCE 8 (bases 18619 to 19770)
AUTHORS Robinson,A.C., Keenan,D.J., Sweeney,J. and Donachie,W.D.
TITLE Further Evidence for Overlapping Transcriptional Units in an *Escherichia coli* Cell Envelope-Cell Division Gene Cluster: DNA Sequence and Transcriptional Organization of the *ddl ftsQ* Region
J. Bacteriol. 167, 809-817 (1986)
MEDLINE 86304170
REFERENCE 9 (bases 22964 to 25011)
AUTHORS Beall,B. and Lutkenhaus,J.
TITLE Sequence analysis, transcriptional organization, and insertional mutagenesis of the *envA* gene of *Escherichia coli*
J. Bacteriol. 169, 5408-5415 (1987)
MEDLINE 88058745
REFERENCE 10 (bases 27412 to 28277)
AUTHORS Akiyama,M., Horituchi,T. and Sekiguchi,M.
TITLE Molecular cloning and nucleotide sequence of the *murT* mutator of *Escherichia coli* that causes A:T to C:G transversion
Mol. Gen. Genet. 206, 9-16 (1987)
MEDLINE 87201091
REFERENCE 11 (bases 843 to 1812)
AUTHORS Henikoff,S., Hauglin,G.W., Calvo,J.M. and Wallace,J.C.
TITLE A large family of bacterial activator proteins
Proc. Natl. Acad. Sci. U.S.A. 85, 6602-6606 (1988)
MEDLINE 88320486
REFERENCE 12 (bases 20513 to 21772)
AUTHORS Robinson,A.C., Beggs,K.J. and Donachie,W.D.
TITLE Mapping and characterization of mutants of the *Escherichia coli* cell division gene, *ftsA*
Mol. Microbiol. 2, 581-588 (1988)
MEDLINE 89039246
REFERENCE 13 (bases 23989 to 27799)
AUTHORS Schmidt,M., Rollo,E., Grodberg,I. and Oliver,D.
TITLE Nucleotide sequence of *seca* gene and *seca*(ts) mutations preventing protein export in *Escherichia coli*
J. Bacteriol. 170, 3404-3414 (1988)
MEDLINE 88298644
REFERENCE 14 (bases 11142 to 12634)
AUTHORS Parquet,C., Fluorel,B., Mengin-Lecreulx,D. and Van Heijenoort,J.
TITLE Nucleotide sequence of the *murF* gene encoding the UDP-MurNac-pentapeptide synthetase of *Escherichia coli*
Nucleic Acids Res. 17, 5379-5379 (1989)
- MEDLINE 89345095
REFERENCE 15 (bases 14743 to 16239)
AUTHORS Ikeda,M., Sato,T., Wachl,M., Jung,H.K., Ishino,F., Kobayashi,M. and Matsunashi,M.
TITLE Structural similarity among *Escherichia coli* *ftsW* and *RodA* proteins and *Bacillus subtilis* SpoVE protein, which function in cell division, cell elongation, and spore formation, respectively
J. Bacteriol. 171, 6375-6378 (1989)
MEDLINE 90036736
REFERENCE 16 (bases 1 to 28277)
AUTHORS Tao,J.S. and Ishiguro,E.E.
TITLE Nucleotide sequence of the *murE* gene of *Escherichia coli*
Can. J. Microbiol. 35, 1051-1054 (1989)
MEDLINE 90124047
REFERENCE 17 (bases 13392 to 15020)
AUTHORS Mengin-Lecreulx,D. and van Heijenoort,J.
TITLE Nucleotide sequence of the *murD* gene encoding the UDP-MurNac-L-Ala-D-Glu synthetase of *Escherichia coli*
Nucleic Acids Res. 18 (1), 183 (1990)
MEDLINE 90174916
REFERENCE 18 (bases 12423 to 15030)
AUTHORS Ikeda,M., Wachl,M., Ishino,F. and Matsunashi,M.
TITLE Nucleotide sequence involving *murD* and an open reading frame *orf-Y* spacing *murF* and *ftsW* in *Escherichia coli*
Nucleic Acids Res. 18, 1058-1058 (1990)
MEDLINE 90192099
REFERENCE 19 (bases 6088 to 7587)
AUTHORS Gomez,M.J., Fluorel,B., Van Heijenoort,J. and Ayala,J.A.
TITLE Nucleotide sequence of the regulatory region of *pbbp* gene of *Escherichia coli*
Nucleic Acids Res. 18, 2813-2813 (1990)
MEDLINE 90251464
REFERENCE 20 (bases 4274 to 6093)
AUTHORS Leclerc,G., Noel,G. and Drapeau,G.
TITLE Molecular cloning, nucleotide sequence and expression of *shl*, a new gene in the 2-minute region of the genetic map of *Escherichia coli*
J. Bacteriol. 172, 4696-4700 (1990)
MEDLINE 90330585
REFERENCE 21 (bases 16094 to 18886)
AUTHORS Ikeda,M., Wachl,M., Jung,H.K., Ishino,F. and Matsunashi,M.
TITLE Nucleotide sequence involving *murG* and *murC* in the *mra* gene cluster region of *Escherichia coli*
Nucleic Acids Res. 18, 4014-4014 (1990)
MEDLINE 90326550
REFERENCE 22 (bases 16094 to 17806)
AUTHORS Mengin-Lecreulx,D., Texier,L. and van Heijenoort,J.
TITLE Nucleotide sequence of the cell-envelope *murG* gene of *Escherichia coli*
Nucleic Acids Res. 18 (9), 2810 (1990)
MEDLINE 90251461
REFERENCE 23 (bases 1 to 28277)
AUTHORS Michaud,C., Parquet,C., Fluorel,B., Blanot,D. and van Heijenoort,J.
TITLE Revised interpretation of the sequence containing the *murE* gene encoding the UDP-N-acetylmuramyl-tripeptide synthetase of *Escherichia coli*
Biochem. J. 269 (1), 277-278 (1990)
MEDLINE 90328986
REFERENCE 24 (bases 1 to 28277)
AUTHORS Wang,Q. and Calvo,J.M.
TITLE Lrp a global regulatory protein of *Escherichia coli* binds co-operatively to multiple sites and activates transcription of *llyV*
J. Mol. Biol. 229, 306-318 (1993)
MEDLINE 93156044
REFERENCE 25 (bases 1 to 28277)
AUTHORS Wang,Q. and Calvo,J.M.
TITLE Lrp a major regulatory protein in *Escherichia coli* bends DNA and can organize the assembly of a higher-order nucleoprotein structure
EMBO J. 12, 2495-2501 (1993)
MEDLINE 93285120
REFERENCE 26 (bases 4274 to 6093)
AUTHORS Jahreis,K., Postma,P.W. and Lengeler,J.W.
TITLE Nucleotide sequence of the *lly H-fur* gene of *Escherichia coli* K-12

JOURNAL Unpublished
 REFERENCE 27 (bases 1 to 28277)
 AUTHORS Ayala,J.A.
 TITLE Regulation of transcription at the 2-minute region of the genetic map of *Escherichia coli*
 JOURNAL Unpublished
 REFERENCE 28 (bases 1 to 28277)
 AUTHORS Ayala,J.A.
 TITLE Direct Submission
 JOURNAL Submitted (08-JAN-1991) Ayala J.A., Instituto de Biologia Molecular, Centro de Biologia Molecular, Universidad Autonoma, Canto-Blanco 28049, Madrid, Spain
 COMMENT This entry comprises a merged sequence of 28kb of which a portion is the submitter's original work.
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Query Match 100.0%; Score 20; DB 1; Length 28277;
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 Db 24817 GATTACGATATAAATCTC 24798

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 DEFINITION Sequence 2 from Patent WO0149775.
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 VERSION AX191720.1 GI:15209889
 KEYWORDS
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 ORGANISM
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 Escherichia coli.
 Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae; Escherichia.
 REFERENCE
 1 (bases 1 to 28277)
 Iversen,P.L.
 TITLE Antisense antibacterial cell division composition and method
 JOURNAL Patent: WO 0149775-A 2 12-JUL-2001;
 Avi Biopharma, Inc. (US)
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 /organism="Escherichia coli"
 /db_xref="taxon:562"

BASE COUNT 6714 a 6725 c 7993 g 6845 t
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Query Match 100.0%; Score 20; DB 6; Length 28277;
 Best Local Similarity 100.0%; Pred. No. 52;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GATTACGATATAAATCTC 20
 Db 24817 GATTACGATATAAATCTC 24798

RESULT 11

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 LOCUS 111401 bp DNA linear BCT 27-JUN-1997
 DEFINITION E.coli K12 genome, 0-2.4min. region.
 ACCESSION D10483 J01597 J01683 J01706 K01298 K01990 M10420 M10611 M12544
 VERSION V00259 X04711 X54847 X54945 X55034 X56742
 D10483.1 GI:216434
 KEYWORDS
 get; DnaJ.
 SOURCE
 Escherichia coli (strain:K-12) DNA.
 ORGANISM
 Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae; Escherichia.
 REFERENCE
 1 (bases 1 to 111401)
 AUTHORS Mori,H.
 TITLE Direct Submission
 JOURNAL Submitted (18-FEB-1992) Hirotsada Mori, Institute for Virus Research, Kyoto University, Genetics and Molecular Biology; 53 Shogoin Kawara Machi, Sakyo-Ku, Kyoto 606, Japan
 (E-mail:es52985@sakura.kudpc.kyoto-u.ac.jp, Tel:075-751-4042, Fax:075-761-5626)
 2 (sites)
 REFERENCE
 2 (sites)
 AUTHORS Smith,B.R. and Schleif,R.
 TITLE Nucleotide sequence of the L-arabinose regulatory region of *Escherichia coli* K12
 JOURNAL The Journal of biological chemistry. 253 (19), 6931-6933 (1978)
 MEDLINE 79005683
 PUBMED 357433
 REFERENCE
 3 (sites)
 AUTHORS Ohtsubo,H. and Ohtsubo,E.
 TITLE Nucleotide sequence of an insertion element, IS1
 JOURNAL Proceedings of the National Academy of Sciences of the United States of America. 75 (2), 615-619 (1978)
 MEDLINE 78137003
 PUBMED 273224
 REFERENCE
 4 (sites)
 AUTHORS Greenfield,L., Boone,T. and Wilcox,G.
 TITLE DNA sequence of the araBAD promoter in *Escherichia coli* B/r
 JOURNAL Proceedings of the National Academy of Sciences of the United States of America. 75 (10), 4724-4728 (1978)
 MEDLINE 79116194
 PUBMED 368797
 REFERENCE
 5 (sites)
 AUTHORS Johnsrud,L.
 TITLE DNA sequence of the transposable element IS1
 JOURNAL Molecular & general genetics : MGG. 169 (2), 213-218 (1979)
 MEDLINE 79177885
 PUBMED 375010
 REFERENCE
 6 (sites)
 AUTHORS Smith,D.R. and Calvo,J.M.
 TITLE Nucleotide sequence of the E coli gene coding for dihydrofolate reductase
 JOURNAL Nucleic acids research. 8 (10), 2255-2274 (1980)
 MEDLINE 81053692
 PUBMED 6159575
 REFERENCE
 7 (sites)
 AUTHORS Miyada,C.G., Horwitz,A.H., Cass,L.G., Tinko,J. and Wilcox,G.
 TITLE DNA sequence of the araC regulatory gene from *Escherichia coli* B/r
 JOURNAL Nucleic acids research. 8 (22), 5267-5274 (1980)
 MEDLINE 81124262
 PUBMED 7008027
 REFERENCE
 8 (sites)
 AUTHORS Ogden,S., Hagerter,D., Stoner,C.M., Kolodrubetz,D. and Schleif,R.
 TITLE The *Escherichia coli* L-arabinose operon: binding sites of the regulatory proteins and a mechanism of positive and negative regulation
 JOURNAL Proceedings of the National Academy of Sciences of the United States of America. 77 (6), 3346-3350 (1980)
 MEDLINE 81013881
 PUBMED 6251457
 REFERENCE
 9 (sites)
 AUTHORS Katinka,M., Cossart,P., Sibillil,L., Saint-Gilons,I., Chalvignac,M.A., Le Bras,G., Cohen,G.N. and Yaniv,M.
 TITLE Nucleotide sequence of the *thra* gene of *Escherichia coli*
 JOURNAL Proceedings of the National Academy of Sciences of the United

MEDLINE 81077247
PUBMED 7003595
REFERENCE 10 (sites)
AUTHORS Mackie, G.A.
TITLE Nucleotide sequence of the gene for ribosomal protein S20 and its flanking regions
JOURNAL The Journal of biological chemistry. 256 (15), 8177-8182 (1981)
MEDLINE 81264207
PUBMED 6267039
REFERENCE 11 (sites)
AUTHORS Cossart, P., Kallinka, M. and Yaniv, M.
TITLE Nucleotide sequence of the thrB gene of E. coli, and its two adjacent regions: the thrAB and thrBC junctions
JOURNAL Nucleic acids research. 9 (2), 339-347 (1981)
MEDLINE 81150470
PUBMED 6259626
REFERENCE 12 (sites)
AUTHORS Lee, N.L., Gielow, W.O. and Wallace, R.G.
TITLE Mechanism of araC autoregulation and the domains of two overlapping promoters, Pc and PBAD, in the L-arabinose regulatory region of Escherichia coli
JOURNAL Proceedings of the National Academy of Sciences of the United States of America. 78 (2), 752-756 (1981)
MEDLINE 81199399
PUBMED 6262769
REFERENCE 13 (sites)
AUTHORS Stoner, C.M. and Schleif, R.
TITLE Is the amino acid but not the nucleotide sequence of the Escherichia coli araC gene conserved?
JOURNAL Journal of molecular biology. 154 (4), 649-652 (1982)
MEDLINE 82216830
PUBMED 6283093
REFERENCE 14 (sites)
AUTHORS Gilson, E., Nikaido, H. and Hofnung, M.
TITLE Sequence of the malK gene in E. coli K12
JOURNAL Nucleic acids research. 10 (22), 7449-7458 (1982)
MEDLINE 83116968
PUBMED 6296778
REFERENCE 15 (sites)
AUTHORS Parsot, C., Cossart, P., Saint-Girons, I. and Cohen, G.N.
TITLE Nucleotide sequence of thrC and of the transcription termination region of the threonine operon in Escherichia coli K12
JOURNAL Nucleic acids research. 11 (21), 7331-7345 (1983)
MEDLINE 84089770
PUBMED 6316258
REFERENCE 16 (sites)
AUTHORS Bouvier, J., Richard, C., Richard, F., Patte, J.C. and Stragier, P.
TITLE Nucleotide sequence and expression of the Escherichia coli darp gene
JOURNAL The Journal of biological chemistry. 259 (23), 14829-14834 (1984)
MEDLINE 85054974
PUBMED 6094578
REFERENCE 17 (sites)
AUTHORS Bardwell, J.C. and Craig, E.A.
TITLE Major heat shock gene of Drosophila and the Escherichia coli heat-inducible dnaK gene are homologous
JOURNAL Proceedings of the National Academy of Sciences of the United States of America. 81 (3), 848-852 (1984)
MEDLINE 84144800
PUBMED 6322174
REFERENCE 18 (sites)
AUTHORS Innis, M.A., Tokunaga, M., Williams, M.E., Lorange, J.M., Chang, S.Y., Chang, S. and Wu, H.C.
TITLE Nucleotide sequence of the Escherichia coli proliptoprotein signal peptidase (lsp) gene
JOURNAL Proceedings of the National Academy of Sciences of the United States of America. 81 (12), 3708-3712 (1984)
MEDLINE 84222028
PUBMED 6374664
REFERENCE 19 (sites)
AUTHORS Bouvier, J., Patte, J.C. and Stragier, P.
TITLE Multiple regulatory signals in the control region of the

JOURNAL Escherichia coli carAB operon
MEDLINE 84248073
PUBMED 6377309
REFERENCE 20 (sites)
AUTHORS Chong, P., Hui, I., Loo, T. and Gillam, S.
TITLE Structural analysis of a new GC-specific insertion element IS186
JOURNAL FEBS letters. 192 (1), 47-52 (1985)
MEDLINE 86030702
PUBMED 2996940
REFERENCE 21 (sites)
AUTHORS Kamio, Y., Lin, C.K., Regue, M. and Wu, H.C.
TITLE Characterization of the lles-lsp operon in Escherichia coli. Identification of an open reading frame upstream of the lles gene and potential promoter(s) for the lles-lsp operon
JOURNAL The Journal of biological chemistry. 260 (9), 5616-5620 (1985)
MEDLINE 85182715
PUBMED 2985604
REFERENCE 22 (sites)
AUTHORS Friedberg, D., Rosenthal, E.R., Jones, J.W. and Calvo, J.M.
TITLE Characterization of the 3' end of the leucine operon of Salmonella typhimurium
JOURNAL Molecular & general genetics. 199 (3), 486-494 (1985)
MEDLINE 85295470
PUBMED 2993799
REFERENCE 23 (sites)
AUTHORS Cowling, D.W., Bardwell, J.C., Craig, E.A., Woolford, C., Hendrix, R.W. and Gross, C.A.
TITLE Consensus sequence for Escherichia coli heat shock gene promoters
JOURNAL Proceedings of the National Academy of Sciences of the United States of America. 82 (9), 2679-2683 (1985)
MEDLINE 85190560
PUBMED 3887408
REFERENCE 24 (sites)
AUTHORS Sekiguchi, T., Ortega-Cesena, J., Nosoh, Y., Ohashi, S., Tsuda, K. and Kanaya, S.
TITLE DNA and amino-acid sequences of 3-isopropylmalate dehydrogenase of Bacillus coagulans. Comparison with the enzymes of Saccharomyces cerevisiae and Thermus thermophilus
JOURNAL Biochim. Biophys. Acta. 867, 36-44 (1986)
MEDLINE 87163495
PUBMED 3549454
REFERENCE 25 (sites)
AUTHORS Lee, N., Gielow, W., Martin, R., Hamilton, E. and Fowler, A.
TITLE The organization of the arabid operon of Escherichia coli
JOURNAL Gene. 47 (2-3), 231-244 (1986)
MEDLINE 86111849
PUBMED 3003084
REFERENCE 26 (sites)
AUTHORS Ohki, M., Tamura, F., Nishimura, S. and Uchida, H.
TITLE Nucleotide sequence of the Escherichia coli dnaJ gene and purification of the gene product
JOURNAL The Journal of biological chemistry. 261 (4), 1778-1781 (1986)
MEDLINE 86111849
PUBMED 3003084
REFERENCE 27 (sites)
AUTHORS

Query Match 100.0%; Score 20; DB 1; Length 111401;
Best Local Similarity 100.0%; Pred. No. 40;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTAGCATTAATAATCTC 20
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Db 107941 GATTAGCATTAATAATCTC 107922

RESULT 12
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LOCUS
DEFINITION
AC055113 Homo sapiens chromosome 12 clone CTD-2021H9, WORKING DRAFT
SEQUENCE, 19 unordered pieces.
AC055113
AC055113.2 GI:9838029
VERSION
KEYWORDS
HTG: HTGS_PHASE1; HTGS_DRAFT.

SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 155976)
AUTHORS Waterston,R.H.
TITLE The sequence of Homo sapiens clone
JOURNAL 2 (bases 1 to 155976)
REFERENCE Waterston,R.H.
AUTHORS Direct Submission
TITLE Submitted (17-APR-2000) Genome Sequencing Center, Washington
JOURNAL University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
On Aug 17, 2000 this sequence version replaced gi:7579846.
COMMENT ----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
----- Project Information -----
Center project name: H_MS2021H09
----- Summary Statistics -----
Sequencing vector: MJ3: 100%
Sequencing vector: plasmid: 0%
Chemistry: Dye-terminator Big Dye; 0% of reads
Chemistry: Dye-terminator Big Dye; 0% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 14393 bases at least Q40
Consensus quality: 146482 bases at least Q30
Consensus quality: 147905 bases at least Q20
Insert size: 18000; agarose-fp
Insert size: 154176; sum-of-ctlgis
Quality coverage: 5.12 in Q20 bases; agarose-fp
Quality coverage: 5.01 in Q20 bases; sum-of-ctlgis

* NOTE: This is a 'working draft' sequence. It currently
* consists of 19 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 3520: contig of 3520 bp in length
* 3521 3620: gap of unknown length
* 3621 9066: contig of 5446 bp in length
* 9067 9166: gap of unknown length
* 9167 13195: contig of 4029 bp in length
* 13196 13295: gap of unknown length
* 13296 17854: contig of 4559 bp in length
* 17855 17954: gap of unknown length
* 17955 26904: contig of 8950 bp in length
* 26905 27004: gap of unknown length
* 27005 35606: contig of 8602 bp in length
* 35607 35706: gap of unknown length
* 35707 43003: contig of 7297 bp in length
* 43004 43103: gap of unknown length
* 43104 52582: contig of 9479 bp in length
* 52583 52682: gap of unknown length
* 52683 62116: contig of 9434 bp in length
* 62117 62216: gap of unknown length
* 62217 75369: contig of 13153 bp in length
* 75370 75469: gap of unknown length
* 75470 88516: contig of 13047 bp in length
* 88517 88616: gap of unknown length
* 88617 104061: contig of 15445 bp in length
* 104062 104161: gap of unknown length
* 104162 119796: contig of 15635 bp in length
* 119797 119896: gap of unknown length
* 119897 141280: contig of 21384 bp in length
* 141281 141380: gap of unknown length
* 141381 142833: contig of 1453 bp in length

FEATURES
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/db_xref="taxon:9606"
/chromosome="12"
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1. 3520
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3621. 9066
/note="assembly_name:Contig11"
9167. 13195
/note="assembly_name:Contig12"
13296. 17854
/note="assembly_name:Contig13"
17955. 26904
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104162. 119796
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141381. 142833
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145183. 148566
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152419. 155976
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155976. 155976
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1802 others
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Best Local Similarity 100.0%; Pred. No. 98;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2 ATTACATTAATAATCTC 20
DB 94819 ATTACATTAATAATCTC 94837
RESULT 13
AC084877/c 179212 bp DNA linear PRI 05-JUN-2001
LOCUS Homo sapiens 12q BAC Rpl1-158011 (Roswell Park Cancer Institute
DEFINITION Human BAC Library) complete sequence.
ACCESSION AC084877
VERSION AC084877.18 GI:14277166
KEYWORDS HTG.
SOURCE human.

ORGANISM	Homo sapiens
1	

Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

REFERENCE AUTHORS

1 (bases 1 to 179212)
Muzny,D.M., Adams,C., Adio-Oduola,B., Ali-osman,F.R., Allen

AUTHORS

[illegible]

Overlapping clones are noted at the beginning and end of the Features listing.

Features listing.

ANNOTATION OF FEATURES:

STSSs are identified using epcC (genome Res. 7:541-550) searches of a local database that includes entries from dbSTS, GDB, and local mapping efforts.

Repeats are identified using RepeatMasker (A. Smit and P. Green, unpublished.) for Human and Mouse sequences.

Genes and Region of sequence similarity are identified by BLAST (Nuc. Acids Res. 25:3389-3402) similarity (expect < 1e-30) to the EST and cDNA sequences. Genes demonstrate at least two exons flanked by consensus splice sites that maintained sequence continuity across the splice junctions. Sequences that are not identical matches are annotated as similar.

SEQUENCING READ COVERAGE: Sequencing is completed to a minimum standard of double strand coverage with a minimum of 2 clones and 2 reads with no ambiguities or 2 chemistries with a minimum of 2 clones and 3 reads with no ambiguities. If the sequence quality for a region does not meet this standard, it will be indicated in the annotation as Low Coverage.

QUALITY OF INDIVIDUAL BASES: this sequence meets stringent quality standards - estimated error rate less than 1 per 10,000 bases. Reports of lowest quality individual bases and measures of base quality are listed below. Description of the metrics can be found at URL: <http://gc.bcm.tmc.edu:8088/quality.info/genbank.annotation.html>.

QUALSTAT-REPORT-----

Summary Statistics

Contig length:	131446
Phrap values in estimate:	131252
Average error rate (BCM-Phrap estimate):	5.08653e-05
Fraction of Phrap values less than 40 :	0.00239996
Number of consensus changing edits:	6
Number of N's in consensus :	0

----- Consensus changing edits

Position	ORF1ginal+Context	ORF1ginal+Context
82935r	tttctcttgt (t)aaagatctct	tttctcttgt (t)aaanattmnc
82962	ctttttaa (g)atctctcag	ctttttaa (n)atmcttcaag
82965	tgttaagat (c)ctctcagag	tgttaagat (n)incttcaagag
82966	gttaagatc (t)ctctcagagc	gttaaanah (n)cttcaagag
82972	gattcttca (g)agctcagaa	gattmctca (n)agctcagaa
82985	gtaccacaa (t)cataccaca	gtattcaca (n)cataccaca

----- Distribution of quality < 40 Bases

# bases	Phrap Value Range
200	5
180	10
160	15
140	20
120	25
100	30
80	35
60	40
40	45
20	50
0	55

FEATURES

Version: 1.01 qxfo.
Location/Qualifiers

CLONE LENGTH: This sequence does not necessarily represent the entire insert of this clone. Overlapping regions of clones are only sequenced and submitted once, so the sequence for the remainder of the insert may be found in the record for the adjacent clones.

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repeat_region
1299..1616
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2804..2898
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3417..3449
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3827..3900
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Query Match 95.0%; Score 19; DB 9; Length 179212;
Best Local Similarity 100.0%; Pred. No. 96;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 ATTAGCATATAAATCTC 20
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Db 10996 ATTAGCATATAAATCTC 10978

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RESULT 14
LOCUS AE008700/c 22286 bp DNA linear BCT 25-OCT-2001
DEFINITION *Salmonella typhimurium* LT2, section 8 of 224 of the complete genome.
ACCESSION AE008700 AE006468
VERSION AE008700.1 GI:16418628
KEYWORDS
SOURCE *Salmonella typhimurium* LT2.
ORGANISM *Salmonella typhimurium* LT2.
Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae; *Salmonella*.
REFERENCE 1 (bases 1 to 22286)
McCelland, M., Sanderson, K.E., Spieth, J., Clifton, S.W., Latreille, P., Courtney, U., Portolillo, S., Ali, J., Dante, M., Du, F., Hou, S., Layman, D., Leonard, S., Nguyen, C., Scott, K., Holmes, A.,

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TITLE
JOURNAL
PUBMED
REFERENCE
AUTHORS
TITLE
JOURNAL

2 (bases 1 to 22286)
The Salmonella typhimurium Genome Sequencing Project.
Direct Submission
Submitted (29-MAR-2001) Genome Sequencing Center, Department of
Genetics, Washington University School of Medicine, 4444 Forest
Park Boulevard, St. Louis, MO 63108, USA
Supported by NIH grant 5U 01 AI43283

COMMENT
Coding sequences below are predicted from manually evaluated
computer analysis, using similarity information and the programs;
GLIMMER: http://www.tigr.org/softlab/glimmer/glimmer.html and
Genemark; http://opal.biology.gatech.edu/Genemark/
EC numbers were kindly provided by Junko Yabuzaki and the Kyoto
Encyclopedia of Genes and Genomes; http://www.genome.ad.jp/kegg/,
and Pedro Romero and Peter Karp at EcoCyc;
http://ecocyc.PangeaSystems.com/ecocyc/
The analyses of ribosome binding sites and promoter binding sites
were kindly provided by Heladia Salgado, Julio Collado-Vides and
ReguionDB;
http://kinich.cifn.unam.mx:8850/db/reguiondb\_intro.frameset
This sequence was finished as follows unless otherwise noted: all
regions were double stranded, sequenced with an alternate
chemistries or covered by high quality data (i.e., phred quality >=
30); an attempt was made to resolve all sequencing problems, such
as compressions and repeats; all regions were covered by sequence
from more than one ml3 subclone.

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Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GATTACGATTAATAAATCTC 20
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Db 7650 GATTAGCATTAATAAAGTCTC 7631

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LOCUS Homo sapiens chromosome 15 clone CTD-2297L20 map 15q21.1, complete
DEFINITION sequence.
AC036102 AC036102.8 GI:16152267
VERSION
KEYWORDS
SOURCE
ORGANISM human.
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
AUTHORS
1 (bases 1 to 127603)
Rowen, L., Madan, A., Qin, S., Baradarani, L., Birditt, B., Bloom, S.,
Burke, J., Dors, M., Fleetwood, P., Kaur, A., Madan, A., Nesbitt, R.,
Pate, D., and Hood, L.
2 (bases 1 to 127603)
Rowen, L., Madan, A., Qin, S., Abbasi, N., Baradarani, L., Birditt, B.,
Bloom, S., Dors, M., Dickhoff, R., Fleetwood, P., Harrison, G.,
James, R., Kaur, A., Madan, A., Owen, M. P., Ratcliffe, A., Shaffer, T.
and Hood, L.

TITLE Direct Submission
JOURNAL Submitted (07-APR-2000) Multimegabase Sequencing Center, University of Washington, PO BOX 357730, Seattle, WA 98195, USA
REFERENCE 3 (bases 1 to 127603)
AUTHORS Rowen, L., Madan, A., Qin, S., Baradarani, L., Birditt, B., Bloom, S., Burke, J., Dots, M., Fleetwood, P., Kaur, A., Madan, A., Nesbitt, R., Pale, D., and Hood, L.
TITLE Direct Submission
JOURNAL Submitted (16-OCT-2001) Multimegabase Sequencing Center, Institute for Systems Biology, 4225 Roosevelt Way NE, Suite 200, Seattle, WA 98105, USA
COMMENT On Oct 16, 2001 this sequence version replaced gi:14318379.
 ----- Genome Center
 Center: Multimegabase Sequencing Center
 Center code: UWMSC
 Web site: http://chroma.mbt.washington.edu/msg_www
 Contact: leetowensystemsbiology.org
 ----- Summary Statistics
 Sequencing vector: pUC18; L08752
 Chemistry: Dye-terminator Big Dye; 90% of reads
 Chemistry: Dye-primer Big Dye; 10% of reads
 Assembly program: Phrap; version 0.990399
 Note: data from AC013452 [Drafting center UWMSC] and AC022306 [Drafting center UWMSC] were added for finishing.
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 DB 61460 GAGTAGCATATATAATCTC 61441

Search completed: October 6, 2002, 16:01:31
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